leaders; families (including families participating in MIECHV-funded home visiting services and those with shared experiences); community members, including community-based program administrators and community service providers, including home visitors.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating, and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS [1]

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Community Interview Protocol	60	1	60	1.50	90
Family and Community Focus Group Guide	240	1	240	2.00	480
Community and Home Visitor Survey Instrument	500	1	500	0.75	375
Program Data	15	1	15	2.00	30
Total	815		815		975

[1] There may be variation in the number of study participants and home visiting programs in each community (*e.g.,* some selected communities may have fewer home visitors). The total burden hours presented here provide information assuming the maximum number of respondents in each community.

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2022–05635 Filed 3–16–22; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0302]

Agency Information Collection Request. 60-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before May 16, 2022.

ADDRESSES: Submit your comments to *Sherrette.Funn@hhs.gov* or by calling (202) 795–7714.

FOR FURTHER INFORMATION CONTACT: When submitting comments or requesting information, please include the document identifier 0990–0302 and project title for reference, to Sherrette A. Funn, email: *Sherrette.Funn@hhs.gov,* or call (202) 795–7714 the Reports Clearance Officer.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection: Medical Reserve Corps Unit Profile and Reports.

Type of Collection: Revision.

OMB No.: 0990–0302.

Abstract: Medical Reserve Corps Units are currently located in 748 communities across the United States and represent a resource of over 300,000 volunteers. In order to continue to support MRC units, detailed information about the MRC units, including unit/user demographics, contact information, volunteer numbers and information about non-emergency and emergency unit activities is needed by the MRC Program. MRC Unit Leaders are asked to update this information on the MRC website at least quarterly and to participate in a technical assistance assessment using the Capability Assessment and Factors for Success at least annually. This collection informs resources and tools developed as part of national programing and helps to identify trends and target technical assistance to support MRC units' preparedness to respond to disasters in their communities. The MRC unit data collection has been refined to eliminate duplication and streamline data collection tools.

ANNUALIZED BURDEN HOUR TABLE

Forms (if necessary)	Respondents (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
Unit Profile	MRC Unit Leader	748	4	15/60	748
Capability Assessment	MRC Unit Leader	748	1	30/60	374
Factors for Success	MRC Unit Leader	748	1	30/60	374

ANNUALIZED BURDEN HOUR TABLE—Continued

Forms (if necessary)	Respondents (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
Unit Activity Reporting	MRC Unit Leader	748	4	15/60	748
Total			10		2,244

Sherrette A. Funn,

Paperwork Reduction Act Reports Clearance Officer, Office of the Secretary.

[FR Doc. 2022–05612 Filed 3–16–22; 8:45 am]

BILLING CODE 4150–47–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Findings of Research Misconduct

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: Findings of research misconduct have been made against Shuo Chen, Ph.D. (Respondent), formerly a postdoctoral researcher, Department of Physics, University of California, Berkeley (UCB). Respondent engaged in research misconduct in research reported in a grant application submitted for U.S. Public Health Service (PHS) funds, specifically National Institute of Neurological Disorders and Stroke (NINDS), National Institutes of Health (NIH), grant application K99 NS116562–01. The administrative actions, including supervision for a period of one (1) year, were implemented beginning on February 28, 2022, and are detailed below.

FOR FURTHER INFORMATION CONTACT: Wanda K. Jones, Dr.P.H., Acting Director, Office of Research Integrity, 1101 Wootton Parkway, Suite 240, Rockville, MD 20852, (240) 453–8200.

SUPPLEMENTARY INFORMATION: Notice is hereby given that the Office of Research Integrity (ORI) has taken final action in the following case:

Shuo Chen, Ph.D., University of California, Berkeley: Based on the report of an investigation conducted by UCB and additional analysis conducted by ORI in its oversight review, ORI found that Dr. Shuo Chen, formerly a postdoctoral researcher, Department of Physics, UCB, engaged in research misconduct in research reported in a grant application submitted for PHS funds, specifically NINDS, NIH, grant application K99 NS116562–01.

ORI found that Respondent engaged in research misconduct by intentionally, knowingly, and/or recklessly falsifying data and methods by altering, reusing, and relabeling source two-photon microscopy and electrophysiological data to represent images of mouse hippocampal neurons in the following grant application:

- K99 NS116562–01, ''Investigation into network dynamics of hippocampal replay sequences by ultrafast voltage imaging,'' submitted to NINDS, NIH, on June 25, 2019.

ORI found that Respondent intentionally, knowingly, and/or recklessly falsified two-photon microscopy and in vivo electrophysiological activity images, figure legends, and text descriptions of hippocampal neurons from a mouse running on a treadmill in a head-fixed virtual reality (VR) set up. Specifically:

- Respondent reused an image of visual cortex neurons to represent fluorescence calcium imaging of hippocampal neurons in Figure 6d and its associated text and figure legend of K99 NS116562–01.
- Respondent reused in vivo electrophysiological data from control mice of spatial receptive fields for all recorded place cells during linear track exploration sessions from Supplemental Figure 1b from *Nat Neurosci.* 2018 Jul;21(7):996–1003 (doi: 10.1038/s41593–018–0163–8) to represent several sessions of two-photon hippocampal calcium imaging of progressive place fields, obtained from multiple mice running on a treadmill in a head-fixed VR set up, in Figure 6e and its associated text and figure legend of K99 NS116562–01.

Respondent neither admits nor denies ORI's findings of research misconduct. The parties entered into a Voluntary Settlement Agreement (Agreement) to conclude this matter without further expenditure of time, finances, or other resources. The settlement is not an admission of liability on the part of the Respondent.

Respondent voluntarily agreed to the following:

(1) Respondent will have his research supervised for a period of one (1) year beginning on February 28, 2022 (the ''Supervision Period''). Prior to the submission of an application for PHS support for a research project on which Respondent's participation is proposed and prior to Respondent's participation in any capacity in PHS-supported research, Respondent will submit a plan for supervision of Respondent's duties to ORI for approval. The supervision plan must be designed to ensure the integrity of Respondent's research. Respondent will not participate in any PHS-supported research until such a supervision plan is approved by ORI. Respondent will comply with the agreed-upon supervision plan.

(2) The requirements for Respondent's supervision plan are as follows:

i. A committee of 2–3 senior faculty members at the institution who are familiar with Respondent's field of research, but not including Respondent's supervisor or collaborators, will provide oversight and guidance during the Supervision Period. The committee will review primary data from Respondent's laboratory on a quarterly basis and submit a report to ORI at six (6) month intervals setting forth the committee meeting dates and Respondent's compliance with appropriate research standards and confirming the integrity of Respondent's research.

ii. The committee will conduct an advance review of each application for PHS funds, or report, manuscript, or abstract involving PHS-supported research in which Respondent is involved. The review will include a discussion with Respondent of the primary data represented in those documents and will include a certification to ORI that the data presented in the proposed application, report, manuscript, or abstract is supported by the research record.

(3) During the Supervision Period, Respondent will ensure that any institution employing him submits, in conjunction with each application for PHS funds, or report, manuscript, or abstract involving PHS-supported research in which Respondent is involved, a certification to ORI that the data provided by Respondent are based on actual experiments or are otherwise legitimately derived and that the data, procedures, and methodology are accurately reported in the application, report, manuscript, or abstract.